IN THE CLAIMS:

11. (Amended) A method as in claim 1, wherein said pyrimidine nucleoside analog is selected from the group consisting of the 5-fluorouracil (5-FU), 5-FU prodrugs including [tegafur] Tegafur and [5'-deoxyfluorouridine, fluorouridine, 2'-deoxyfluorouridine] 5'-deoxy-5-fluorouridine, 5-fluorouridine, 2'-deoxy-5-fluorouridine, prodrug derivatives of [fluorouridine or 2'-deoxyfluorouridine] 5-fluorouridine or 2'-deoxy-5-fluorouridine, fluorocytosine, trifluoromethyl-2'-deoxyuridine, arabinosyl cytosine, prodrugs of arabinosyl cytosine, cyclocytidine, 5-aza-2'-deoxycytidine, arabinosyl 5-azacytosine, 6-azacytidine, N-phosphonoacetyl-L-aspartic acid (PALA), pyrazofurin, 6-azauridine, azaribine, thymidine, and 3-deazauridine.